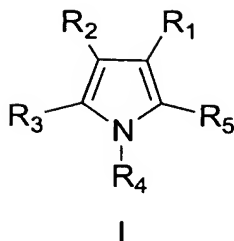


CLAIMS:

1. A compound of Formula I



or a pharmaceutically acceptable salt thereof, wherein

10 R_1 and R_2 are independently selected from optionally substituted aryl and optionally substituted heteroaryl;

R_3 is selected from hydrogen, optionally substituted alkyl, $-N=CR'''$, $-C(O)R'$, $-C(O)NR'R''$, $-NR'R''$, optionally substituted aryl, optionally substituted heteroaryl, and optionally substituted heterocycle, wherein R' and R'' are independently selected from hydrogen, optionally substituted alkyl, optionally substituted aryl, and optionally substituted heterocycle;

20 R_4 is selected from hydrogen, optionally substituted alkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted heterocycle, and $-SiR'''R''''R'''''$ wherein R''' , R'''' , and R''''' are each an independent straight chain or branched C_{1-5} alkyl, or R_3 , R_4 and the $-C-N-$ to which R_3 and R_4 are connected together form an optionally substituted 5- or 6-membered ring;

25 R_5 is selected from optionally substituted alkyl, $-C(O)OR'$, $-C(O)NR'R''$, $C(O)NHNHC(O)R_6$, $-SO_2NR'R''$, $-C(O)R'$, $-NR'R''$, nitrile, nitro, halo, and optionally substituted heterocycle, or R_4 , R_5 and the $-C-N-$ to which R_4 and R_5 are connected together form an optionally substituted 5- or 6-membered ring;

30

R₆ is selected from H, alkyl, optionally substituted aryl; and




with the provisos that

5

(1) R₁ and R₂ are not both optionally substituted phenyl;

(2) if either R₁ or R₂ is optionally substituted phenyl or 3-thienyl, and the

10

other is unsubstituted ,  or , then R₃ is not hydrogen, unsubstituted alkyl, -(CH₂)₃OH, -(CH₂)₃PH, -(CH₂)₃OMs, or -(CH₂)₂N(CH₂)₂O(CH₂)₂, and R₅ is not unsubstituted alkyl, -(CH₂)₃OH, -(CH₂)₃PH, -(CH₂)₃OMs, or -(CH₂)₂N(CH₂)₂O(CH₂)₂; and

(3) R₄ doesn't form a fused ring with both R₃ and R₅.

15

2. The compound of Claim 1 wherein R₁ is substituted with a group selected from hydrogen, amino, alkyl substituted amino, aryl substituted amino, hydroxy, methoxy, phenyl ether, S-alkyl, halogen, trifluoromethyl, and nitro.

20

3. The compound of Claim 1 wherein R₂ is substituted with a group selected from hydrogen, amino, alkyl substituted amino, aryl substituted amino, hydroxy, methoxy, phenyl ether, S-alkyl, halogen, trifluoromethyl, and nitro.

4. The compound of Claim 3 wherein R₂ is heteroaryl having 1-3 N.

25

5. The compound of Claim 1 wherein R₃ is selected from hydrogen, alkyl, aryl, heteroaryl, heterocycle, and -NR'R'', wherein R' and R'' are independently selected from hydrogen, alkyl, aryl, and heterocycle.

30

6. The compound of Claim 1 wherein R₄ is hydrogen or alkyl.

7. The compound of Claim 6 wherein R₄ is hydrogen or methyl.

8. The compound of Claim 1 wherein R_5 is selected from alkyl, $-C(O)OR'$, $-C(O)NR'R''$, nitrile, and heterocycle.

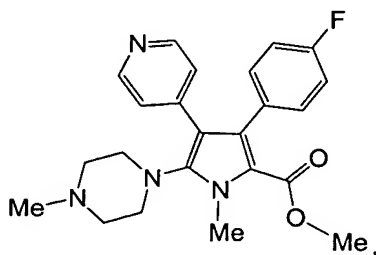
5 9. The compound of Claim 8 wherein R_5 is selected from is selected from $-(CH_2)_nOR'$, $-(CH_2)_nNR'R''$, $-(CH_2)_nCOOR'$, $-(CH_2)_nCONR'R''$, $-NHCOR'$, and ester isosteres.

10 10. The compound of Claim 9 wherein R_5 is selected from oxadiazole, 1,2,4-triazole, 1,2,4-triazol-3-ol, isoxazol-3-ol, imidazolidine-2,4-dione, 4H-[1,2,4]thiadiazol-5-one, oxazole, and [1,3,4]oxadiazole.

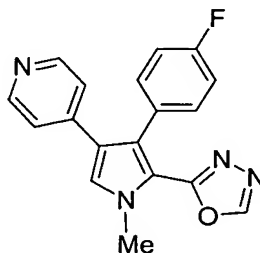
11. The compound of Claim 10 wherein R_5 is 4H-[1,2,4]oxadiazole-5-thione or 4H-[1,2,4]oxadiazol-5-one.

15

12. The compound of claim 1 having the structure

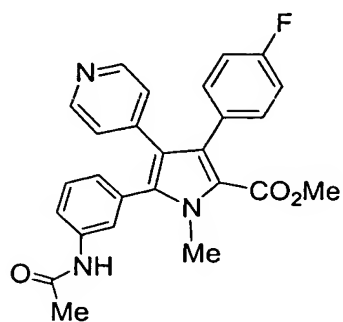


13. The compound of claim 1 having the structure

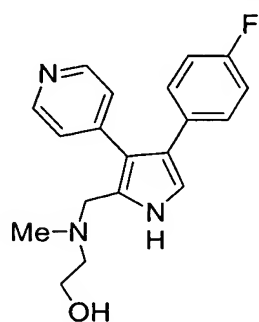


20

14. The compound of claim 1 having the structure

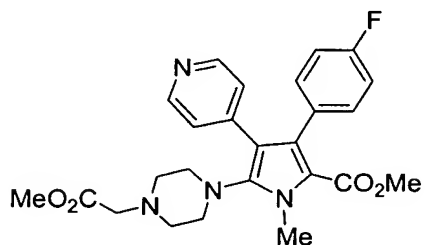


15. The compound of claim 1 having the structure



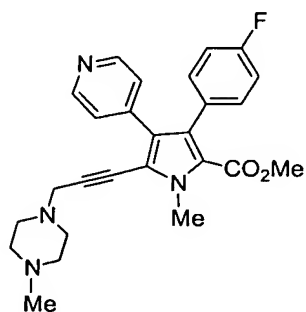
5

16. The compound of claim 1 having the structure

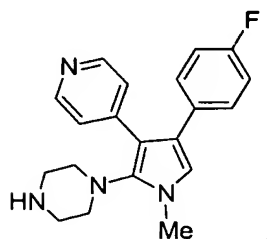


10

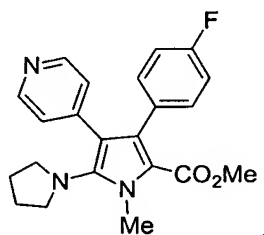
17. The compound of claim 1 having the structure



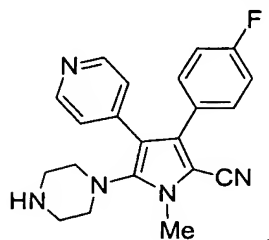
18. The compound of claim 1 having the structure



5 19. The compound of claim 1 having the structure

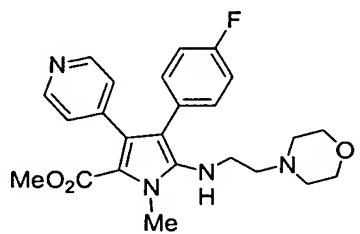


20. The compound of claim 1 having the structure



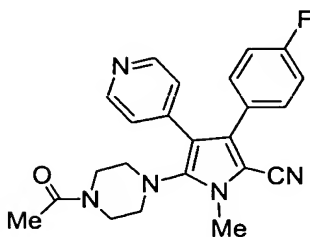
10

21. The compound of claim 1 having the structure

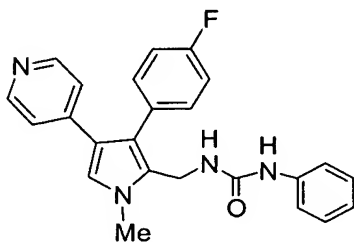


15

22. The compound of claim 1 having the structure



23. The compound of claim 1 having the structure



5

24. A pharmaceutical composition comprising the compound of claim 1 and a pharmaceutically acceptable carrier.

10 25. A method of treating a subject having a disorder ameliorated by reducing TNF- α production and/or p38 activity in appropriate cells, which comprises administering to the subject a therapeutically effective dose of the pharmaceutical composition of claim 24.

15 26. The method of claim 25, wherein the disorder is an inflammatory disorder.

27. The method of claim 25, wherein the disorder is selected from the group consisting of rheumatoid arthritis, osteoporosis, osteoarthritis, allergic inflammation, periodontal disorder, inflammatory bowel disorder, septic shock,
 20 insulin-dependent diabetes mellitus, non-insulin-dependent diabetes, cachexia, pulmonary fibrosis, myasthenia gravis, Crohn's disease, hepatitis, primary biliary cirrhosis, acute pancreatitis, allograft rejection, glioblastoma, alopecia areata, psoriasis, ischemia, congestive heart failure, restenosis, atherosclerosis, systemic lupus erythematosus, nephritis, Guillain-Barre Syndrome, viral
 25 myocarditis, HIV replication, T-cell depletion in HIV infection, cognitive deficits

induced by neuronal inflammation, multiple sclerosis, stroke, neuropathic pain, HIV dementia and Alzheimer's disease.

28. The method of claim 27, wherein the disorder is rheumatoid arthritis.

5

29. A method of preventing an inflammatory response in a subject, comprising administering to the subject a prophylactically effective amount of the pharmaceutical composition of claim 24 either preceding or subsequent to an event anticipated to cause the inflammatory response in the subject.

10

30. The method of claim 29, wherein the event is selected from the group consisting of an insect sting and an animal bite.

15